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(54) Title: ANTIBODIES FOR USE IN ANTILYMPHOCYTE ANTIBODY THERAPY (57) Abstract <p>The use of antibodies against human alpha tumour necrosis factor in antilymphocyte antibody therapy is described. The anti-human alpha tumour necrosis factor antibodies may be used to prevent or treat shock-related conditions arising from antilymphocyte antibody therapy. Also described are compositions containing anti-human alpha tumour necrosis factor antibodies and antilymphocyte antibodies, as well as recombinant anti-human alpha tumour necrosis factor antibodies.</p>		

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ANTIBODIES FOR USE IN ANTILYMPHOCYTE ANTIBODY THERAPY

Field of the Invention

This invention relates to antibodies against human alpha (α)-tumour necrosis factor for use in antilymphocyte antibody therapy, to therapeutic compositions containing an antibody against human α -tumour necrosis factor and an antilymphocyte antibody, and to recombinant antibodies against human α -tumour necrosis factor.

Background to the Invention

Antilymphocyte antibody therapy has become a recognised clinical technique for use when it is desired to supplement or modify the normal immune response in humans. Thus, for example, the intravenous administration of polyclonal rabbit or horse antilymphocyte antibodies is an effective treatment for acute kidney allograft rejection. More recently, monoclonal antilymphocyte antibodies, such as Orthoclone OKT3, have been used for this purpose.

Despite the effectiveness of antilymphocyte antibody therapy, treatment has been hindered in many cases by the occurrence of shock-related side effects, making it necessary to temporarily discontinue the antibody infusion. Side effects include fever and chills, arthralgias, nausea and vomiting, tachycardia, angina pectoris, dyspnoea due to bronchospasm, and, in the case of OKT3 infusion, pulmonary oedema. In addition to the discomfort these side effects can cause, the severity of some of the effects precludes the use of antilymphocyte antibody therapy in some patients, for example people with pulmonary or cardiac disease.

We have now found that patients who are undergoing antilymphocyte antibody therapy, and who are also experiencing shock-related side effects, have surprisingly high plasma levels of α -tumour necrosis factor (α -TNF). We have used this to develop means to control shock-related conditions arising from antilymphocyte antibody therapy.

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Summary of the Invention

Thus, according to one aspect of the invention we provide an antibody against human α -tumour necrosis factor for use in the prevention or treatment of shock-related conditions arising from antilymphocyte antibody therapy.

The term antilymphocyte antibody therapy as used herein is especially to be understood to mean the use of an antilymphocyte antibody in the prophylaxis or treatment of immunoregulatory disorders in which rejection of self or non-self tissue occurs, for example in autoimmune diseases such as thyroiditis or rheumatoid arthritis, or, in particular, in a rejection episode following an organ or tissue transplant.

The shock-related conditions which may arise from antilymphocyte antibody therapy and which may be prevented or treated according to the present invention may be any physiological conditions which are generally associated with a degree of circulatory collapse. Particular conditions include, for example, cardiac conditions such as tachycardia and angina, e.g. angina pectoris; pulmonary conditions such as bronchospasm; musculo-skeletal conditions, for example joint pains such as arthralgia; and general metabolic disorders such as oedema, e.g. pulmonary oedema, and abnormal body temperatures.

The antibodies against human α -tumour necrosis factor (hereinafter referred to as anti-TNF α antibodies) for use according to the invention may in general belong to any immunoglobulin class. Thus for example the anti-TNF α antibody may be an immunoglobulin G or immunoglobulin M antibody.

The anti-TNF α antibody may be of animal, for example mammalian origin and may be for example of murine, rat or human origin. The antibody may be a whole immunoglobulin, or a fragment thereof, for example a fragment derived by proteolytic cleavage of a whole antibody, such as F(ab')₂, Fab' or Fab fragments, or fragments obtained by recombinant DNA techniques, for example Fv fragments (as

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The anti-TNF α antibody may be polyspecific but is preferably monospecific for human α -TNF. The antibodies may be polyclonal or monoclonal antibodies. Particularly useful antibodies for use according to the invention include recombinant anti-TNF α antibodies and fragments thereof, i.e. anti-TNF α antibodies or fragments which have been produced using recombinant DNA techniques. Such recombinant anti-TNF α antibodies are novel compounds and form another aspect of the invention.

Especially useful recombinant antibodies include, (1) those having an antigen binding site at least part of which is derived from a different antibody, for example those in which the hypervariable or complementarity determining regions of one antibody have been grafted into the variable framework regions of a second, different antibody (as described in European Patent Specification No. 239400); (2) recombinant antibodies or fragments wherein non-Fv sequences have been substituted by non-Fv sequences from other, different antibodies (as described in European Patent Specifications Nos. 171496, 173494 and 194276); or (3) recombinant antibodies or fragments possessing substantially the structure of a natural immunoglobulin but wherein the hinge region has a different number of cysteine residues from that found in the natural immunoglobulin, or wherein one or more cysteine residues in a surface pocket of the recombinant antibody or fragment is in the place of another amino acid residue present in the natural immunoglobulin (as described in International patent Applications Nos. PCT/GB 88/00730 and PCT/GB 88/00729 respectively).

The anti-TNF α antibodies may be prepared using well-known immunological techniques employing α -TNF as antigen. Thus, for example, any suitable host may be injected with α -TNF and the serum collected to yield the desired polyclonal anti-TNF α antibody after appropriate purification and/or concentration, (for example by affinity chromatography using immobilised α -TNF as the affinity medium). Alternatively, splenocytes or lymphocytes may be recovered from the α -TNF-injected host and immortalised using for example the method of Kohler et al., Eur. J. Immunol. 6, 511, (1976), the

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resulting cells being segregated to obtain a single genetic line producing monoclonal anti-TNF α antibodies in accordance with conventional practice. Antibody fragments may be produced using conventional techniques, for example by enzymatic digestion of whole antibodies e.g. with pepsin [Parham, J. Immunol., 131, 2895, (1983)] or papain [Lamoyi and Nisonoff, J. Immunol. Meth., 56, 235, (1983)]. Where it is desired to produce recombinant anti-TNF α antibodies these may be produced using for example the general methods described in the above-mentioned patent specifications.

In order to prevent or treat shock-related conditions arising from antilymphocyte antibody therapy, anti-TNF α antibodies may in general be administered in an appropriate form and amount at any suitable time before or during the therapy and, where necessary, after the therapy has finished. The anti-TNF α antibodies may be administered separately or together with antilymphocyte antibodies.

Thus according to a further aspect of the invention we provide a two component system, each component for use in association with one another in antilymphocyte antibody therapy, said system comprising (1) a first component that is an anti-TNF α antibody, and (2) a second component that is an antilymphocyte antibody.

Where the anti-TNF α and antilymphocyte antibodies are administered separately, each may be formulated according to conventional practice. Thus, according to a further aspect of the invention, we provide a pharmaceutical composition comprising an anti-TNF α antibody together with one or more pharmaceutically acceptable carriers, excipients or diluents for use in the prevention or treatment of shock conditions arising from antilymphocyte antibody therapy.

The compositions according to this aspect of the invention may contain other active ingredients.

Where it is desired to administer the anti-TNF α and antilymphocyte antibody together, these may be conveniently

formulated in the same composition. Thus according to a further aspect of the invention we provide a pharmaceutical composition which comprises an anti-TNF α antibody and an antilymphocyte antibody in admixture with one or more pharmaceutically acceptable carriers, excipients or diluents.

In yet a further aspect of the invention we provide a pharmaceutical composition which comprises an anti-TNF α antibody, and an antilymphocyte antibody in admixture with one or more pharmaceutically acceptable carriers, excipients or diluents, for use in antilymphocyte antibody therapy.

In a still further aspect of the invention, we provide a method for the manufacture of a pharmaceutical composition for use in the prevention or treatment of shock conditions arising from antilymphocyte antibody therapy which comprises admixing an anti-TNF α antibody and one or more pharmaceutically acceptable carriers, excipients or diluents.

In the compositions according to the invention the antilymphocyte antibody may be for example a polyclonal rabbit or horse antilymphocyte antibody, or a monoclonal antilymphocyte antibody such as Orthoclone OKT3. Such antibodies are readily available from known sources.

The compositions may take any suitable form for administration, and in particular will be in a form suitable for parenteral administration e.g. by injection or infusion, for example by bolus injection or continuous infusion. Compositions for injection or infusion may take such forms as suspensions, solutions or emulsions of the antibody in oily or aqueous vehicles, and, may contain formulatory agents such as suspending, stabilising and/or dispensing agents. Alternatively, the composition may be in a dry form, for reconstitution before use with an appropriate sterile liquid.

The dose at which the anti-TNF α antibody will be administered will depend on the nature of the antilymphocyte antibody therapy in use

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and whether the anti-TNF α antibody is being used prophylactically or to treat an existing shock-related condition arising from antilymphocyte antibody therapy. Thus, for example, the anti-TNF α antibody may be administered by infusion to a 70kg man, usually at a total dose in the range 20 - 500mg, over for example 2 - 3 days or for as long as the shock condition arising from the antilymphocyte antibody therapy persists.

Where the anti-TNF α antibody is infused concurrently with an antilymphocyte antibody, the antilymphocyte antibody may be infused at generally accepted doses depending on the antibody in use, for example with Orthoclone OKT3 at a dose of 5mg/day for 10 -14 days in a 70kg man.

In a further aspect of the invention we provide a method of treatment of a human subject suffering from an immunoregulatory disorder which comprises administering effective amounts of an anti-TNF α antibody and an antilymphocyte antibody in association to the human subject.

Immunoregulatory disorders include, for example, those described above.

In another aspect of the invention we provide a method of treatment of a human subject suffering from a shock condition arising from antilymphocyte antibody therapy which comprises administering an effective amount of an effective amount of an anti-TNF α antibody to the human subject.

Description of Specific Embodiments

Patients

Informed consent was obtained from seven consecutive kidney transplant patients, treated for acute allograft rejection with antithymocyte globulin (ATG) at the Maastricht University Hospital, to take serial blood samples for plasma tumour necrosis factor (TNF)

measurement during ATG treatment. ATG treatment consisted of intravenous administration of 100-300 mg ATG together with 50 mg prednisolone in 500 ml saline, infused over a 8-hr period. All patients received 4 mg of the antihistamine chlorpheniraminemaleate orally 1/2 hr before start of the infusion. EDTA-anticoagulated 5 ml blood samples were drawn from an intravenous line inserted in the arm opposite the one in which ATG was infused. Blood samples were immediately centrifuged and aliquots of plasma were stored at -70°C for measurement of TNF concentration. During treatment patient temperature and blood pressure were monitored and the occurrence of other side effects of the treatment was registered.

Enzyme-linked Immunosorbent Assay (ELISA) for TNF

Plasma TNF concentrations were determined with a TNF-specific ELISA using monoclonal and polyclonal anti-TNF antibodies. Monoclonal antibodies were derived from anti-TNF secreting hybridomas obtained by a standard cell fusion procedure. Two different monoclonal antibodies were coated overnight at 4°C in a flatbottom microtiter plate (Greiner, Nurtingen, FRG) at a final concentration of 5 μg Ig/ml. Plates were blocked with 1% (w/v) bovine serum albumin. Samples to be tested for TNF concentration were added to the wells and incubated for 1 1/2 hrs. at room temperature. A standard titration curve was obtained by making serial dilutions of a known sample of recombinant TNF. The amount of TNF bound to the wells was quantitated by sequential incubation with a 1:2000 dilution of rabbit immune serum and a 1:5000 dilution of peroxidase conjugated goat anti-rabbit antibody (Jackson, West Grove, PA; antibody not crossreactive with human serum proteins), followed by adding substrate (o-phenylenediamine: Sigma, St. Louis, MO) to the wells. The colour reaction was stopped with 0.1 M H_2SO_4 and absorption of light at 495 nm was measured with a microelisa autoreader (Flow, Irvine, UK). The lower detection limit of ELISA ranged between 5-10 pg/ml in plasma. TNF specific values were obtained by subtracting background absorptions due to non-TNF plasma proteins cross-linking in the ELISA, measured in the absence of either the monoclonal anti-TNF antibodies coated on the bottom of the ELISA plate or the second, rabbit anti-TNF antibodies. Plasma

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samples of a large group (n=60) of healthy volunteers were shown to be negative for TNF (i.e. < 5 pg/ml) as determined with our assay.

In vitro experiments

Peripheral blood mononuclear cells (PBMC) were prepared by buoyant density centrifugation on Lymphoprep (Nyegaard, Oslo, Norway) of buffy coats of donor blood. Monocytes were obtained by depleting PBMC of T-lymphocytes through rosetting with sheep red blood cells and centrifugation on Lymphoprep after which the interface layer cells were left to adhere to plastic for 1/2 hr at 37°C. After extensive washing, to remove non-adherent cells, the cells were shown to consist of more than 95% of esterase-positive cells. PBMC and monocytes were cultured in RPMI 1640 (Gibco, Paisley, Scotland) supplemented with 10% heat-inactivated (56°C, 20 min.) foetal calf serum (Boehringer, Mannheim, FRG) and 100 IU/ml penicillin and 100 ug/ml streptomycin (Flow, Irvine, UK). PBMC and monocytes were cultured in cluster wells (24-well plates, Greiner); In each well 2.5×10^8 PBMC and monocytes obtained after adherence of 2.5×10^8 T lymphocyte depleted PBMC per well were cultured in 600 ul medium. To these cultures different concentrations of ATG and of the anti-CD3 monoclonal antibody OKT3 (ascites derived from an OKT3-secreting hybridoma, purchased from the ATCC, Rockville, MD) were added and at different times samples of cell-free supernatant were harvested to measure TNF concentration. Lymphocyte proliferation was determined by measuring the incorporation of ^3H -thymidine. 0.5 uCi ^3H -thymidine (Amersham, UK; specific activity 5.0 Ci/mmol) was added to a sample of 100ul of the cell cultures transferred into round-bottomed microtiter plates (Greiner) and after a 4 hour culture period the radioactivity incorporated in the cells was measured with a liquid scintillation counter.

TNF Levels in patients treated with ATG

All 7 patients treated for kidney graft rejection with ATG showed elevated plasma TNF levels, starting about 1 hour and reaching peak levels between 2-3 hrs after the start of the ATG infusion. Plasma

TNF returned to zero level at about 6 hrs after the beginning of therapy. All patients developed a rise in body temperature shortly after plasma TNF had reached peak values. Body temperature normalized on the average 6 hr after TNF disappeared from the circulation. In most of the patients a slight rise in blood pressure accompanied with an increased heart rate was observed during the ATG infusion. All patients experienced mild to severe side effects, which coincided in time with the period of elevated plasma TNF levels, and which made it necessary to temporarily discontinue the ATG-infusion in several patients. In all patients symptoms waned shortly after TNF plasma levels had returned to normal. All patients experienced a feeling of great fatigue at the end of the therapy. Table I gives the relevant data on all 7 patients. Maximum plasma TNF concentrations in these patients ranged between 111 and 731 pg/ml on the first day of treatment. On the second day of treatment, when side effects of the ATG infusion were considerably less, maximum TNF levels were much lower, i.e. between 0 and 55 pg/ml.

TNF-production by peripheral blood mononuclear cells in vitro

In vitro experiments were performed in order to elucidate the cell source and mechanism of production of TNF in these patients. ATG or OKT3 was added at different concentrations to cultures of peripheral blood mononuclear cells (PBMC) and after a 6 hour culture period, TNF concentration in the supernatant was determined. Both ATG and OKT3 induced in a dose-related manner the release of TNF by PBMC in vitro. OKT3, which is a T-cell mitogen, also induced T-lymphocyte proliferation. However, ATG induced TNF secretion was not accompanied by T-lymphocyte proliferation, indicating that proliferation of T-lymphocytes is not a necessary prerequisite for TNF production in these cultures.

The kinetics of the ATG- and OKT3-induced release of TNF by PBMC was compatible with the rapid release of TNF in the circulation of the ATG-treated patients.

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Table 1. Plasma TNF Concentrations And Other Relevant Data of ATG-treated Patients

Patient*	treatment day	treatment	highest TNF conc. (pg/ml)	highest temp (°C)	RR** (mm Hg)	symptoms
1 m 33yr	1	300mg ATG	560	39.4	+50/+5	chills, vomiting
	2	200mg ATG	55	37.9	0/0	none
2 f 54yr	1	100mg ATG	179	39.5	+10/0	chills, dyspnoea
3 f 35yr	1	200mg ATG	111	39.0	+5/+5	chills
	2	200mg ATG	24	38.0	0/0	none
4 m 49yr	1	200mg ATG	125	38.4	-30/-20	chills, dyspnoea, vomiting
	2	200mg ATG	11	37.2	0/0	none
5 f 61yr	1	200mg ATG	138	38.2	+15/0	none
6 f 54yr	1	200mg ATG	731	40.4	+10/-10	chills, dyspnoea, headache
	2	100mg ATG	0	37.4	0/0	none
7 m 43yr	1	200mg ATG	593	39.5	+15/-10	chills, diarrhea, headache

* m = male; f = female

**Expressed as difference between systolic/diastolic blood pressure measured before start of the ATG infusion and at the time of the highest plasma TNF concentration

CLAIMS

1. A pharmaceutical composition which comprises an antibody against human α -tumour necrosis factor and an antilymphocyte antibody in admixture with one or more pharmaceutically acceptable carriers, excipients or diluents.
2. A pharmaceutical composition according to Claim 1 wherein the antilymphocyte antibody is a monoclonal antilymphocyte antibody.
3. A pharmaceutical composition according to Claim 2 wherein the monoclonal antilymphocyte antibody is Orthoclone OKT3.
4. A pharmaceutical composition according to any of Claims 1, 2, or 3, for use in antilymphocyte antibody therapy.
5. A two component system, each component for use in association with one another in antilymphocyte antibody therapy, said system comprising (1) a first component that is an antibody against human α -tumour necrosis factor, and (2) a second component that is an antilymphocyte antibody.
6. An antibody against human α -tumour necrosis factor for use in the prevention or treatment of shock-related conditions arising from antilymphocyte antibody therapy.
7. A pharmaceutical composition comprising an antibody against human α -tumour necrosis factor together with one or more pharmaceutically acceptable carriers, excipients or diluents for use in the prevention or treatment of shock conditions arising from antilymphocyte antibody therapy.
8. An antibody against human α -tumour necrosis factor characterised in that the antibody is prepared by a recombinant DNA technique.

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9. A method of treatment of a human subject suffering from an immunoregulatory disorder which comprises administering effective amounts of an anti-TNF α antibody and an antilymphocyte antibody in association to the human subject.
10. A method of treatment of a human subject suffering from a shock condition arising from antilymphocyte antibody therapy which comprises administering an effective amount of an antibody against human α -tumour necrosis factor to the human subject.

INTERNATIONAL SEARCH REPORT

International Application No

PCT/GB 89/00254

I. CLASSIFICATION OF SUBJECT MATTER (if several classification symbols apply, indicate all) ⁴ According to International Patent Classification (IPC) or to both National Classification and IPC IPC ⁴ : A 61 K 39/395											
II. FIELDS SEARCHED <div style="text-align: center;">Minimum Documentation Searched ⁷</div> <table style="width: 100%; border: none;"> <tr> <td style="width: 50%; border: none;">Classification System </td> <td style="width: 50%; border: none;">Classification Symbols</td> </tr> <tr> <td style="border: none;">IPC⁴ </td> <td style="border: none;">C 12 P</td> </tr> </table> <div style="text-align: center; padding-top: 10px;">Documentation Searched other than Minimum Documentation to the Extent that such Documents are Included in the Fields Searched ⁸</div>			Classification System	Classification Symbols	IPC ⁴	C 12 P					
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III. DOCUMENTS CONSIDERED TO BE RELEVANT ⁹ <table border="1" style="width: 100%; border-collapse: collapse;"> <thead> <tr> <th style="width: 10%;">Category ¹⁰</th> <th style="width: 70%;">Citation of Document, ¹¹ with indication, where appropriate, of the relevant passages ¹²</th> <th style="width: 20%;">Relevant to Claim No. ¹³</th> </tr> </thead> <tbody> <tr> <td style="text-align: center; vertical-align: top;">A</td> <td>EP, A, 0218868 (NEW YORK BLOOD CENTER) 22 April 1987</td> <td></td> </tr> <tr> <td style="text-align: center; vertical-align: top;">A</td> <td>Chemical Abstracts, vol. 106, 1987, (Columbus, Ohio, US), C.P. Robinson: "Muromonab-CD-3 (Orthoclone OKT3) - a review", see page 511, abstract 154301t, & Med. Actual. 1986 22(12), 603-9 -----</td> <td></td> </tr> </tbody> </table>			Category ¹⁰	Citation of Document, ¹¹ with indication, where appropriate, of the relevant passages ¹²	Relevant to Claim No. ¹³	A	EP, A, 0218868 (NEW YORK BLOOD CENTER) 22 April 1987		A	Chemical Abstracts, vol. 106, 1987, (Columbus, Ohio, US), C.P. Robinson: "Muromonab-CD-3 (Orthoclone OKT3) - a review", see page 511, abstract 154301t, & Med. Actual. 1986 22(12), 603-9 -----	
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<div style="display: flex; justify-content: space-between;"> <div style="width: 45%;"> <p>¹⁰ Special categories of cited documents: ¹⁰</p> <p>"A" document defining the general state of the art which is not considered to be of particular relevance</p> <p>"E" earlier document but published on or after the international filing date</p> <p>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</p> <p>"O" document referring to an oral disclosure, use, exhibition or other means</p> <p>"P" document published prior to the international filing date but later than the priority date claimed</p> </div> <div style="width: 45%;"> <p>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</p> <p>"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step</p> <p>"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art</p> <p>"&" document member of the same patent family</p> </div> </div>											
IV. CERTIFICATION <table style="width: 100%; border: none;"> <tr> <td style="width: 50%; border: none;">Date of the Actual Completion of the International Search 6th July 1989</td> <td style="width: 50%; border: none;">Date of Mailing of this International Search Report 27. 7. 89</td> </tr> <tr> <td style="border: none;">International Searching Authority EUROPEAN PATENT OFFICE</td> <td style="border: none;">Signature of Authorized Officer A. J. VAN DER PUTTEN</td> </tr> </table>			Date of the Actual Completion of the International Search 6th July 1989	Date of Mailing of this International Search Report 27. 7. 89	International Searching Authority EUROPEAN PATENT OFFICE	Signature of Authorized Officer A. J. VAN DER PUTTEN					
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